

Form PTO-1449		Docket Number 295002005900		Application Number 09/175,017	
INFORMATION DISCLOSURE CITATION IN AN APPLICATION (Use several sheets if necessary)		Applicant BOIME et al			
		Filing Date 19 October 1998		Group Art Unit 1645 1647	
U.S. PATENT DOCUMENTS					
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Examiner Initials	Ref. No.	Date	Document No.	Country	Class Subclass Translation YES NO
OTHER DOCUMENTS (including author, title, Date, Pertinent Pages, Etc.)					
Examiner Initials	Ref. No.	Title			
J V	1.	LaPolt, P.S. "Enhanced Stimulation of Follicle Maturation and Ovulatory Potential by Long Acting Follicle-Stimulating Hormone Agonists with Extended Carboxyl-Terminal Peptides", <i>Endocrinology</i> (1992) 131:2514-2520.			
	2.	Fares, F.A.. "Design of a Long Acting Follitropin Agonist by Fusing the C-terminal Sequence of the Chorionic Gonadotropin β subunit to the Follitropin" <i>Proc Natl Acad Sci USA</i> (1992) 89:4304-4308.			
	3.	Lapthorn, A.J. "'Crystal Structure of Human Chorionic Gonadotropin" <i>Nature</i> (1994) 369:455-461.			
	4.	Wu, H. "Structure of Human Chorionic Gonadotropin at 2.6 A Resolution from MAD analysis of the Selenomethionyl Protein" <i>Structure</i> (1994) 2:545-558.			
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	6.	Dayhoff, M. "A Model of Evolutionary Change in Proteins" <i>Atlas of Protein Sequences and Structure</i> (1972) 5:89-99.			
	7.	Chen, F. "The Carboxy-Terminal Region of the Glycoprotein Hormone α -Subunit: Contributions to Receptor Binding and Signaling in Human Chorionic Gonadotropin" <i>Molec Endocrinol</i> (1992) 6:914-919.			
	8.	Yoo, J. "Conversion of Lysine 91 to Methionine or Glutamic Acid in Human Choriogonadotropin α Results in the Loss of cAMP Inducibility" <i>J Biol Chem</i> (1991) 266:17741-17743.			
	9.	Puett, D. "Delineation of Subunit and Receptor Contact Sites by Site-Directed Mutagenesis of hCG β " <i>Glycoprotein Hormones</i> .			
	10.	Lusbader, J.W., <i>EDS, Springer Verlag</i> (New York) (1994) 122-134.			
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Paper #4

Form PTO-1449

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Docket Number 295002005900

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Applicant

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Group Art Unit ~~1648~~ 1647

OTHER DOCUMENTS

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V	21.	Sachais "Molecular Basis for the Species Selectivity of the Substance P Antagonist CP-96, 345" <i>Biol Chem</i> (1993) 268:2319.

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